## **CLAIMS**

- 1. A method of crystallizing a macrolide from a macrolide starting material comprising the steps of:
- a) combining a macrolide starting material, a polar solvent, a hydrocarbon solvent,
  and water, whereby at least two phases are formed, at least one of which is a water-rich phase, and wherein the pH of the water-rich phase is at least about 7,
  - b) maintaining the combination at for at least 1 hour, whereby a macrolide-rich phase is formed from which the macrolide crystallizes.
- 10 2. The method of claim 1 further comprising the step of isolating the macrolide that crystallizes.
  - 3. The method of claim 1 wherein the combination of step b is maintained at a temperature of from about -15°C to about 50°C.

- 4. The method of claim 3 wherein the combination of step b is maintained at a temperature of from about -5°C to about 40°C.
- 5. The method of claim 4 wherein the combination of step b is maintained at a temperature of from about -2°C and about 35°C.
  - 6. The method of claim 1 wherein the combination of step b is maintained for between 48 and 100 hours.
- 7. The method of claim 1 wherein the polar solvent is selected from the group consisting of alcohols, esters, nitriles and ethers.
  - 8. The method of claim 7 wherein the polar solvent is selected from the group consisting of ethyl acetate, acetonitrile, methanol, ethanol, *n*-propanol, *iso*-propanol, *n*-

butanol, iso-butanol, acetone, diisopropyl ether, dimethyl formamide, and dimethyl acetamide.

9. The method of claim 8 wherein the polar solvent is ethyl acetate.

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- 10. The method of claim 1 wherein the hydrocarbon solvent is selected from the group consisting of n-hexane, n-heptane, octane, iso-octane cyclohexane, methylcyclohexane, benzene, toluene, and xylene.
- 10 11. The method of claim 10 wherein the hydrocarbon solvent is *n*-hexane.
  - 12. The method of claim 1 wherein the pH of the water-rich phase is about 8 or higher.
- 15 13. The method of claim 1 wherein the water comprises a base selected from NaOH, KOH, Ca(OH)<sub>2</sub>, NH<sub>3</sub>, Et<sub>3</sub>N, diethylamine and pyridine.
  - 14. The method of claim 1 wherein the macrolide is selected from the group consisting of tacrolimus, sirolimus, pimecrolimus, and everolimus.
  - 15. A method of crystallizing a macrolide from a macrolide starting material comprising the steps of:
  - a) combining a concentrate residue from whole-broth extraction of macrolide-containing biomatter in a polar solvent with a hydrocarbon solvent, and water, whereby at least two phases are formed, at least one of which is a water-rich phase, and wherein the pH of the water-rich phase is at least about 7,
  - b) maintaining the combination at for at least 1 hour, whereby a macrolide-rich phase is formed from which the macrolide crystallizes.

- 16. The method of claim 15 further comprising the step of isolating the macrolide that crystallizes.
- 17. The method of claim 15 wherein the combination of step b is maintained at a temperature of from about -15°C to about 50°C.
  - 18. The method of claim 17 wherein the combination of step b is maintained at a temperature of from about -5°C to about 40°C.
- 10 19. The method of claim 18 wherein the combination of step b is maintained at a temperature of from about -2°C and about 35°C.
  - 20. method of claim 15 wherein the combination of step b is maintained for between 48 and 100 hours.
  - 21. The method of claim 15 wherein the polar solvent is selected from the group consisting of alcohols, esters, nitriles and ethers.

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- 22. The method of claim 21 wherein the polar solvent is selected from the group consisting of ethyl acetate, acetonitrile, methanol, ethanol, *n*-propanol, *iso*-propanol, *n*-butanol, *iso*-butanol, acetone, diisopropyl ether, dimethyl formamide, and dimethyl acetamide.
  - 23. The method of claim 22 wherein the polar solvent is ethyl acetate.
  - 24. The method of claim 15 wherein the hydrocarbon solvent is selected from the group consisting of *n*-hexane, *n*-heptane, octane, *iso*-octane cyclohexane, methylcyclohexane, benzene, toluene, and xylene.

- 25. The method of claim 24 wherein the hydrocarbon solvent is *n*-hexane.
- 26. The method of claim 15 wherein the pH of the water-rich phase is about 8 or higher.
- 27. The method of claim 15 wherein the water comprises a base selected from NaOH, KOH, Ca(OH)<sub>2</sub>, NH<sub>3</sub>, Et<sub>3</sub>N, diethylamine and pyridine.
- 28. The method of claim 15 wherein the macrolide is selected from the group consisting of tacrolimus, sirolimus, pimecrolimus, and everolimus.

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- 29. A method of crystallizing a macrolide from a macrolide starting material comprising the steps of:
- a) combining, at a temperature of about 20° to about 25°C, macrolide starting

  material, ethyl acetate, *n*-hexane, and a water solution of a base selected from NaOH,

  KOH, Ca(OH)<sub>2</sub>, NH<sub>3</sub>, (C<sub>2</sub>H<sub>5</sub>)<sub>3</sub>N, diethylamine and pyridine whereby at least two phases

  are formed, one of which is a water-rich phase, wherein the pH of the water-rich phase is

  > about 7,
- b) maintaining the combination at a temperature of about 20°C to about 25°C for
   at least 1 hour, whereby a macrolide-rich phase is formed from which macrolide crystallizes,
  - c) maintaining the combination at a temperature of about 0°C to about 20°C for at least 1 hour, and
    - d) recovering the macrolide that crystallizes.
  - 30. The method of claim 29 wherein the macrolide is selected from the group consisting of tacrolimus, sirolimus, pimecrolimus, and everolimus.
- 31. The method of claim 29 wherein the pH of the water-rich phase is about 8 or 30 higher.

- 32. A method of crystallizing a macrolide from a macrolide starting material comprising the steps of:
- a) combining, at a temperature of about 20° to about 25°C, a concentrate residue from whole-broth extraction of macrolide-containing biomatter in ethyl acetate, *n*-hexane, and a water solution of a base selected from NaOH, KOH, Ca(OH)<sub>2</sub>, NH<sub>3</sub>, (C<sub>2</sub>H<sub>5</sub>)<sub>3</sub>N, diethylamine and pyridine whereby at least two phases are formed, one of which is a water-rich phase, wherein the pH of the water-rich phase is > about 7,
  - b) maintaining the combination at a temperature of about 20°C to about 25°C for at least 1 hour, whereby a macrolide-rich phase is formed from which macrolide crystallizes,
    - c) maintaining the combination at a temperature of about 0°C to about 20°C for at least 1 hour, and
      - d) recovering the macrolide that crystallizes.

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- 33. The method of claim 32 wherein the macrolide is selected from the group consisting of tacrolimus, sirolimus, pimecrolimus, and everolimus.
- 34. The method of claim 32 wherein the pH of the water-rich phase is about 8 or 20 higher.
  - 35. In a method for crystallizing a macrolide from a macrolide starting material, the step of combining the macrolide starting material, a polar solvent, a hydrocarbon solvent, and water, whereby at least two phases are formed, at least one of which is water rich, wherein the pH of the water-rich phase is at least about 7.
  - 36. In a method for crystallizing a macrolide from a concentrate residue from whole-broth extraction of macrolide-containing biomatter in a polar solvent, the step of combining the macrolide concentrate in the polar solvent, a hydrocarbon solvent, and water, whereby at least two phases are formed, at least one of which is water rich, wherein the pH of the water-rich phase is at least about 7.